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Title

Inventors

Takeshi FUJITA,

Takashi IRIE,

Kazunori OKANO.

PREPARATION METHOD OF NUCLEIC ACID SAMPLE FOR RARE
EXPRESSED GENES AND ANALYZING METHOD USING THE PREPARED
NUCLEIC ACID SAMPLES THEREBY

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BACKGROUND OF THE INVENTION

1. Field of the Invention

10 The present invention relates to a preparation
method of nucleic acid sample, in particular to a
preparation method of nucleic acid sample suitable for
analysis of rare expressed genes in the sample, to nucleic
acid samples obtained by the preparation method and to an
analyzing method using the prepared nucleic acid samples.

2. Description of the Related Art

15 Constitutive cells of a human body can be classified
into 200 or more species, and each one cell type has subtly
different subtypes. Each of the cells has a common
genome coding for about 10×10^4 genes and several ten
thousands genes are expressed therein according to its
cell type. Investigations on such gene expression have
20 become more important not only for obtaining findings on
functions of individual genes but also for clarifying
biological phenomena. Detailed analyses on
comparatively limited few genes have revealed that plural
genes operate in coordination in majority of the
25 biological phenomena. In the course of the human genome
project, investigations on networks of genes for
macroscopic comprehension of relations between genes and
biological phenomena are to start. Under these

circumstances, demands have been made to provide techniques for further detailed analyses of a multitude of genes.

Recently, there have been developed analyzing
5 methods of gene expression by the comparison between RNA
fingerprintings (e.g., differential display method (Liang,
P., and Pardee, A. B. (1992) Science 257,967-971) and
molecular index method (Kikuya Kato (1995) Nucleic Acids
Res., 23,3685-3690)) and methods using DNA chips,
10 suggesting the possibility of complete analysis of
expressed genes.

Expressed Genes can be roughly classified according
to its expressing amount into three classes, i.e.,
abundant class where a gene is expressed 12,000 copies or
15 more per cell, intermediate class where a gene is expressed
about 300 copies per cell and rare class where a gene is
expressed about 15 copies. In mammals, species of
expressed genes amount to several ten thousands per cell
and most genes belong to the rare class. In other words,
20 expressed genes in a cell comprise enormous species of rare
expressed genes coresident with extremely few species of
abundant expressed genes yet expressed on three or four
orders of magnitude greater than the rare expressed genes
(e.g., Alberts, B., et al. (1989) Molecular Biology of the
25 Cell, 2nd edition, Garland Publishing Inc., NY, USA).

As analyzing methods of rare expressed gene are known
an equalized library (Minoru S. H. Ko (1990) Nucleic Acids
Res., 18,5705-5711), methods based upon multiplex PCR

(polymerase chain reaction) including the differential display method and the molecular index method, and methods using DNA chips. It is known, however, that the equalized library technique would lose information on expression amounts because the amounts of genes are made uniform among gene species; and that in an analysis using multiplex PCR such as in the differential display method a comparative PCR has a strong bias towards abundant expressed genes to result in less sensitivity than a usual PCR and thereby to fail to detect rare expressed genes (David J. Bertioli, et al. (1995) Nucleic Acids Res., 23,4520-4523).

SUMMARY OF THE INVENTION

It is, therefore, an object of the present invention to provide a preparation method of nucleic acid sample suitable for analyses of rare expressed genes, and nucleic acid samples prepared thereby, and to provide an analyzing method of rare expressed genes using the prepared nucleic acid samples.

The present inventors focused that abundant expressed gene (abundant class) in a nucleic acid sample can relatively easily be analyzed and readily removed in a selective manner, as there is a strong bias towards abundant expressed genes in the process of gene expression analysis using the differential display method, and hence the expression analysis of rare expressed genes using a nucleic acid obtained by removing abundant expressed genes in advance from a nucleic acid sample to be analyzed is expected to achieve higher precision. The present

invention has been accomplished on the basis of the above findings.

BRIEF DESCRIPTION OF THE DRAWINGS

These and other features, objects, and advantages of the present invention will become apparent upon a consideration of the following description of the invention when read in conjunction with the drawings, in which:

FIG. 1 graphically illustrates a fundamental procedure of an analyzing method of rare expressed genes according to the invention;

FIGS. 2A through 2G graphically illustrate a portion of the flow of an embodiment of the latter three steps among five steps for obtaining a nucleic acid sample predominantly composed of rare expressed genes from a nucleic acid sample to be analyzed, as shown in FIG. 1;

FIGS. 3A through 3G graphically illustrate a portion of the flow of another embodiment of the latter three steps among five steps for obtaining a nucleic acid sample predominantly composed of rare expressed genes from a nucleic acid sample to be analyzed, as shown in FIG. 1;

FIGS. 4A through 4E graphically illustrate residual following the steps shown in FIGS. 2A through 2G or FIGS. 3A through 3G;

FIGS. 5A through 5G graphically illustrate the flow of a partially modified embodiment of the embodiment shown in FIGS. 2A through 2G;

FIGS. 6A through 6F graphically illustrate a portion

of the flow of a still another embodiment of the latter three steps among five steps for obtaining a nucleic acid sample predominantly composed of rare expressed genes from a nucleic acid sample to be analyzed, as shown in FIGS.

5 2A through 2G;

FIG. 7 is a cross sectional view illustrating an embodiment of an apparatus which is useful for the recovery of nucleic acid samples not immobilized on a probe as in embodiments according to the invention;

10 FIGS. 8A and 8B graphically illustrate, respectively, a sectional view and usage of, and a plane view of another embodiment of an apparatus which is useful for the recovery of nucleic acid sample not hybridized with a probe in the embodiments according to the invention;

15 FIG. 9 graphically illustrates the flow of a further embodiment according to the invention of the steps for obtaining a nucleic acid sample predominantly composed of rare expressed genes from a nucleic acid samples to be analyzed; and

20 FIG. 10 is a photograph illustrating the result of electrophoresis on an acrylamide gel to verify the advantages of the invention.

DESCRIPTION OF THE PREFERRED EMBODIMENT

25 FIG. 1 graphically illustrates fundamental procedures of an analyzing method of rare expressed genes according to the invention. Process 100 is a preparing process of a nucleic acid sample to be analyzed, process 200 is a process for removing one or plural abundant

expressed genes in the nucleic acid sample by adding and hybridizing probe carriers which have complement sequence of the above genes, process 300 is a process for obtaining a nucleic acid sample predominantly composed of rare
5 expressed genes by recovering nucleic acid sample not being hybridized with the probe carriers, and process 400 is a process for analyzing the rare expressed genes. Details of the processes 100 through 300, that is, steps for obtaining a nucleic acid sample predominantly composed
10 of rare expressed genes from the nucleic acid sample to be analyzed are as follows:

- Step 1: a preparation step of a nucleic acid sample from a biological material;
- Step 2: a synthesizing step of probe carriers;
- 15 Step 3: a step of mixing and hybridizing the nucleic acid with the probe carriers;
- Step 4: a step of removing genes hybridized with the probe carriers; and
- Step 5: a step of recovering a nucleic acid sample for
20 rare expressed genes.

In the above steps, the term "nucleic acid sample" means a mixture composed of, for example, mRNAs (messenger RNAs) derived from plural genes extracted from a
biological material to be investigated such as cells,
25 tissues or individuals, total RNA inclusive of mRNAs, or cDNAs (complementary DNAs) synthesized based upon mRNAs. Genes are classified according to the amounts into the abundant, intermediate and rare classes. The term "genes

to be removed" in the invention means genes belonging to the abundant class and some of the intermediate class, and the term "genes to be analyzed" means genes belonging to the other of the intermediate class, and those belonging to the rare class.

Embodiments of the present invention will be described below, but they do never limit the scope of the invention.

In Step 1, any known technique according to a biological material to be subjected can be applied. By taking purification of RNA as nucleic acid sample, an AGPC method (e.g., Chomczynski, P. and Sacchi, N. (1987) Anal. Biochem., 162,156-159) can for example be applied, using a commercially available reagent such as TRIZOL Reagent (GIBCO BRL, Gaithersburg, MD., USA). A first strand cDNA can be synthesized according to the method of Sambrook, J. et al. (Molecular Cloning: A Laboratory Manual, 2nd ed. Cold Spring Harbor Laboratory Press, NY, USA) using a commercially available kit.

In Step 2, a gene to be removed from a nucleic acid sample (hereinafter referred to as "LEES GENE") should be selected. Information on genes which are comparatively abundantly expressed in biological materials can be obtained through, for instance, currently publicly displayed data bases such as BodyMap. BodyMap (<http://www.imcb.osaka-u.ac.jp/bodymap/>) is established by Kousaku Okubo, Kenichi Matsubara et al. (IMCB, Osaka Univ., Japan) and is a database of expressed genes for

individual human tissues. Even when there is no information, such genes can be studied comparatively easily by experiments using, for example, SAGE method (Victor E. Velculescu et al. (1995) Science, 270,484-487). Most of LEES GENES are expected to be so called housekeeping genes which are expressed ubiquitously in most species of cells. If only in the identical species, therefore, most of such selected gene species can be adopted to another biological material by modifying only a portion of the selected genes.

The probe sequence should be designed to meet the following conditions: that a sequence which is complementary to a portion of the sequence of the LEES GENE and is hardly hybridized with other gene should be selected; that probe carriers should not hybridized with each other; that a sequence should be selected which is hardly form intermolecular hydrogen bonds; and that the melting temperature (T_m) between the probe carrier and the target gene should be in a proper range (preferably a range from 40° to 70°C). The configuration, material and other factors of the probe carrier vary from a removing method, and these will be described later.

In Steps 3 through 5, the following three methods can be employed. The first method, second method and third method will be further described with reference to, respectively, FIGS. 2A through 2G, 3A through 3G, 4A through 4E and 5A through 5G; FIGS. 6A through 6F, 7 and 8A through 8B; and FIG. 9. FIGS. 2A through 2G, 3A through

3G, 4A through 4E and 5A through 5G illustrate an example where LEES GENE is one gene, whereas when plural LEES GENES should be removed, plural species of probe carriers which are obtained by designing and synthesizing individual probe carriers independently with respect to individual LEES GENES in a nucleic acid sample are provided and the removal of the LEES GENES can be conducted in turn.

Initially, the first method will now be described in detail.

The embodiment illustrated in FIGS. 2A through 2G are conducted as follows:

FIG. 2A illustrates the state of a provided nucleic acid sample, which contains plural species of rare expressed genes 1 to be remained, and gene 2 to be removed (hereinafter referred to as "LEES GENE 2". As the nucleic acid sample, a nucleic acid sample composed of mRNAs, as well as total RNA, can be employed. Although the figure gives the impression that the LEES GENE 2 is less in amount than the rare expressed genes 1 in the sample, the LEES GENE 2 belongs to the abundant class and, in actuality, occupies relatively larger part than the rare expressed genes in the sample.

FIG. 2B illustrates the step of adding a probe carrier to the provided nucleic acid sample. As the probe carrier 3 is used DNA having a sequence hybridizable with the LEES GENE 2 in the vicinity of the 3' end, or its analogue, such as one composed of PNA (peptide nucleic acid) whose 3' end is modified to inhibit elongation. Any material which has

high specificity to the sequence and is recognizable by an enzyme can be used as a material of the probe carrier 3.

5 The probe carrier 3 used in this embodiment will be briefly described below.

Most of DNA synthesis methods using an automated synthesizer are solid phase synthesis methods in which 3' end nucleoside is bonded to a resin such as aminomethyl-polystyrene or controlled-pore glass (CPG),
10 and after the last nucleotide is bound therewith, the obtained DNA is cut from the resin. When a probe carrier is prepared according to this method, a further modification is required to obtain a probe carrier having a modified 3' end. A DNA synthesis method, which uses
15 a resin-bonded nucleoside having a structure as to prevent a elongation reaction, can avoid re-modification. As example of such a method, there may be mentioned a DNA synthesis method using a resin-bonded nucleoside in which the bonding site between the nucleoside and the resin is
20 changed from 3'- to 2'-position and the hydroxyl group in the 3'-position is changed to a hydrogen atom

The use of the aforementioned method therefore facilitates the synthesis of probe carriers each having modified 3' end corresponding to a variety of LEES GENES.
25 In addition, the application of the method to a commercially available synthesizer can provide probe carriers without the need for special instruments and techniques.

FIG. 2C illustrates the state where the probe carrier 3 is hybridized with the LEES GENE 2 in the provided nucleic acid sample. In this step, the nucleic acid sample and the probe carrier 3 are mixed in, for example, a microtube, and the mixture is allowed to stand under conditions (time, temperature of the reaction mixture, concentration of a salt, tramp materials) suitable for hybridization of the LEES GENE 2 with the probe carrier 3. In addition to a microtube, use can be made of any reactor whose wall rarely adsorbs nucleic acids and whose structure can prevent evaporation of a nucleic acid sample solution, and whose temperature can be controlled externally or internally. The condition of hybridization can be calculated using a software for probe design such as OLIGO (trade name, National Biosciences, Inc., Plymouth, Minn., USA).

FIG. 2D illustrates a treatment of adding Ribonuclease H to the nucleic acid sample in which the probe carrier 3 and the LEES GENE 2 hybridize with each other.

FIG. 2E illustrates the state where the sequence region of the LEES GENE 2 to which the probe carrier 3 is specifically and sufficiently hybridized is digested by the Ribonuclease H. By the digestion, the LEES GENE 2 is digested in the vicinity of 3' end to give a fragment 4. Some of Ribonuclease H have activity in a wide range of thermostability and other hybridization conditions, including *E. coli* Ribonuclease H and *Thermus thermophilus* Ribonuclease H (TOYOBO, Japan). Any of such Ribonuclease

H can be employed depending on the hybridization conditions. In addition to Ribonuclease H, enzymes or compounds that can act and digest specifically nucleic acids hybridized with probes may be employed.

5 FIG. 2F illustrates an inactivation treatment of the Ribonuclease H added to the nucleic acid sample; and

 FIG. 2G illustrates the removal of the probe 3 from the nucleic acid sample by ,for example, treating the sample with a DNase.

10 A nucleic acid sample is thus prepared from which the LEES GENE 2 hybridized with the probe carrier 3 has been removed.

 Next, the embodiment shown in FIGS. 3A through 3G will be illustrated below:

15 The treatment to the LEES GENE as illustrated in FIGS. 2A through 2G can be improved for more efficient removal by using a plurality of probe carriers. FIGS. 3A through 3G demonstrate this technique.

20 FIG. 3A illustrates the state of a provided nucleic acid sample as in FIG. 2A.

 FIG. 3B illustrates, as in FIG. 2B, a treatment of adding probe carriers to the provided nucleic acid sample, whereas in this embodiment, two species of probes having different sequences are employed as the probe carriers 3 and 3'. In this treatment, as the probe carrier 3 is used DNA having a sequence hybridizable with the LEES GENE in the vicinity of the 3' end, or its analogue, such as one composed of a PNA (peptide nucleic acid), whereas as the

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probe carrier 3' is employed DNA having a sequence hybridizable with the LEES GENE at a site away from its 3' end, or its analogue, such as one composed of a PNA (peptide nucleic acid). Any materials which have high
5 specificity to the sequences and are recognizable by an enzyme can advantageously be used as materials of the probe carriers 3 and 3'. The similar probe carriers to those described in FIG. 2B can preferably be employed in this embodiment.

10 FIG. 3C illustrates, as in FIG. 2C, the state where the probe carriers are hybridized with the LEES GENE 2 in the prepared nucleic acid sample. As two species of probe carriers 3 and 3' are employed in this embodiment, there are three possible situations, i.e., with the LEES GENE
15 2, the probe carrier 3 is hybridized; the probe carrier 3' is hybridized; and both the probe carriers 3 and 3' are hybridized.

FIG. 3D illustrates, as in FIG. 2D, the treatment of adding Ribonuclease H to the nucleic acid where the
20 probe carriers 3 and 3' and the LEES GENE 2 hybridize with each other.

FIG. 3E illustrates the state where the sequence regions of the LEES GENE 2 to which either or both of the probe carriers 3 and 3' are specifically and sufficiently
25 hybridized are digested by the Ribonuclease H. As there are three ways of the hybridization of the probe carriers 3 and 3' in this embodiment, there are three ways of the digestion by the added Ribonuclease H. By the digestion,

the LEES GENE 2 is digested in the vicinity of its 3' end to give a fragment 4 as in FIG. 2E. Any of various Ribonuclease H, as well as other enzymes and compounds, as described in FIG. 2E can be employed in the present embodiment.

FIG. 3F illustrates an inactivation treatment of the Ribonuclease H added to the nucleic acid sample; and

FIG. 3G illustrates the removal of the probe 3 and/or 3' from the nucleic acid sample by, for example, treating the sample with a DNase.

As described above, the use of plural probe carriers can provide the hybridization between a LEES GENE and probe carriers with higher reliability and hence provide the removal of the LEES GENE with higher efficiency. The species of the probe carriers is naturally not limited to two species, and three or more species of probes can be employed.

As the probe carriers to be used in the present embodiment, those exemplified in the previous embodiment can be used.

Next, the embodiment shown in FIGS. 4A through 4E will now be illustrated:

In this embodiment, cDNA is synthesized from mRNA as the nucleic acid sample prepared in the manner as in FIGS. 2A through 2G.

FIG. 4A illustrates the state of the prepared nucleic acid sample by the above method. To be more specific, in the sample, rare expressed genes 1 which are plural species

of mRNAs are coresident with the LEES GENE 4 resulted from the digestion of the LEES GENE 2 in the vicinity of its 3' end.

FIG. 4B illustrates a treatment of adding a primer to the nucleic acid sample to synthesis cDNA. In this embodiment, an oligo (dT) primer is used, whereas the primer to be used depends on a technique for expression analysis. By way of illustration, a primer obtained by adding any of A, G or C to the 3' end of an oligo(dT) primer is, for example, used in the differential display method.

FIG. 4C illustrates the state of a proceeding cDNA synthesis reaction. The cDNA synthesis reaction is conducted using a reverse transcriptase (e.g., Super Script II RT, a product of GIBCO BRL, Gaithersburg, MD., USA). The cDNA synthesis reaction for the rare expressed genes 1 proceeds in a normal manner to give cDNA 5, whereas that for the LEES GENE 4 does not proceed and no cDNA 5 is synthesized. This is because priming site of the LEES GENE 4 is cleaved and removed, and the synthesis reaction ceases at the site of cleavage.

FIG. 4D illustrates a treatment of treating the nucleic acid sample with an RNase to remove RNAs from the sample.

FIG. 4E illustrates the state of the thus-prepared nucleic acid sample composed of cDNA 5 containing no LEES GENE 2.

Detailed description of the technique for synthesizing cDNA from mRNA, the nucleic acid sample

obtained in the manner shown in FIGS. 3A through 3G, is omitted herein. In this case, as in FIGS. 4A through 4E, cDNA synthetic reaction ceases at the site that has been digested due to the hybridization with the probe carrier 3 and/or 3' to give a nucleic acid sample composed of cDNA containing no LEES GENE 2.

In the first method mentioned above, the digestion reaction and subsequent treatment steps illustrated in FIGS. 2D through 2G and FIGS. 3D through 3G can be omitted. To this end, for example, a method using a probe carrier (e.g., a ribozyme) having cleaving activity on a hybridized region can be employed. The use of such a carrier to be hybridized with the LEES GENE as shown in FIGS. 2C and 3C naturally results in the state illustrated in FIG. 4A without the procedure of FIG. 2D or FIG. 3D.

The embodiment illustrated in FIGS. 5A through 5E will now be described below:

FIG. 5A illustrates the state of a nucleic acid sample composed of mRNAs to be used in this embodiment. The nucleic acid sample is composed of plural species of rare expressed genes 1 to be remained coresident with the LEES GENE 2, as in FIG. 2A.

FIG. 5B illustrates a treatment of adding probe carrier to the provided nucleic acid sample. As the probe carrier 3, use is made of a probe carrier having a sequence hybridizable with the LEES GENE 2 in the vicinity of its 3' end, being designed to have a T_m (melting temperature) of at least equal to, and ideally higher than that of primer

for cDNA synthesis, and 3' end of the probe is modified to avoid elongation. The probe carrier may be composed of DNA or its analogue such as PNA (peptide nucleic acid) that has high specificity to the sequence and is resistant against the digestion by a reverse transcriptase. The DNA in this case should be modified with, for example, phosphorothioates since DNA as intact is liable to be digested. The 3' end of the probe carrier can be modified by deoxylation or amination, for instance. The probe carrier to be used in this embodiment should preferably one described in relation with FIG. 2B.

FIG. 5C illustrates the state where the probe carrier 3 is hybridized with the LEES GENE 2 in the provided nucleic acid sample. The hybridization can be conducted in a similar manner as in FIG. 2C.

FIG. 5D illustrates a treatment of adding a primer to the nucleic acid sample to synthesize cDNA. In this embodiment, an example using oligo(dT) primer is employed as in FIG. 4B, whereas the synthetic primer and conditions of cDNA synthetic reaction can be modified as in the aforementioned embodiment.

FIG. 5E illustrates the state of a proceeding cDNA synthesis reaction. The cDNA synthesis reaction can be carried out under similar conditions as in FIG. 4C. The synthesis of cDNA 5 from the LEES GENE 2 is inhibited because the probe carrier 3 is hybridized with the LEES GENE 2 in the vicinity of its 3' end.

FIG. 5F illustrates a treatment of removing RNA from

the nucleic acid sample by treating the sample with an RNase.

FIG. 5G illustrates the state of the thus-prepared nucleic acid sample composed of cDNA 5 containing no LEES GENE 2.

In this case, the primer for cDNA synthesis is added (FIG. 5D) after hybridizing the probe carrier with the LEES GENE (FIG. 5C), whereas cDNA synthesis can be conducted after adding the probe carrier 3 and the primer concurrently to the sample and then allowing the reaction condition to meet the hybridization condition. The bond between the probe carrier and LEES GENE is made through hydrogen bonds in this case, whereas it may also be conducted by, for example, providing a photoaffinity-labeled probe carrier 3 and hybridizing it with the LEES GENE 2 and then allowing the LEES GENE and probe carrier to combine with each other through covalent bonds by light irradiation. A similar probe carrier as demonstrated with reference to FIG. 2B can be employed as the probe carrier in this embodiment.

Next, the second method according to the invention will be described in detail with reference to FIGS. 6A through 6F. The second method is characterized by immobilizing a probe on a solid phase. As the nucleic acid sample, any of mRNAs and first strand cDNAs can be employed. The sequence of the probe can be directed to any of regions in the LEES GENE, but when the region to be analyzed in the expression analysis is expected, a probe coding for

the identical region with the target region should preferably be employed. By taking the differential display method as example, the 3' region is analyzed and hence a probe coding for 3' region is to be used. The solid phase for immobilizing the probe can have any configuration, such as in the form of bead, glass plate or microtube. It should immobilize the probe with high density and the adsorption of DNA by its surface should be minimized. The use of beads, in particular, of magnetic beads will be described in this embodiment.

FIG. 6A illustrates a treatment of providing a solution 12 containing a target nucleic acid sample in a microtube 11. The nucleic acid sample may be composed of mRNAs or total RNA, as in described with reference to FIG. 2A. The sample contains LEES GENE 16 with plural rare expressed genes 15 to be remained.

FIG. 6B illustrates a treatment of preparing bead 13 onto which probe carrier 14 being complementary to the LEES GENE 16 is immobilized. As the bead, commercially available beads such as Dynabeads (registered trade mark; DYNAL CORPORATION, Oslo, Norway) and BioMag (Perseptive Biosystems, Bedford, Mass., USA) can be employed. The immobilization can for example be conducted in the following manner: Probe 14 whose 5' end is biotinylated is synthetically obtained, and mixed with beads whose surfaces are coated with streptavidin in a buffer to immobilize the probe onto the beads through avidin-biotin bonds.

FIG. 6C illustrates a treatment of mixing the provided immobilized probe 14 with the nucleic acid sample 12 in the microtube 11, allowing the reaction mixture under conditions suitable for hybridization (the conditions can be referred to those in the first method), and specifically hybridizing the LEES GENE 16 with the probe 14.

Consequently, the LEES GENE 16 is hybridized with the probe 14 and thus captured onto the surfaces of beads, whereas the rare expressed genes 15 to be remained are including in the nucleic acid sample.

FIG. 6D illustrates a treatment of collecting beads cluster 17 to an area of the microtube 11 with the use of a magnet.

FIGS. 6E and 6F illustrate a treatment of remaining the beads cluster 17 in the microtube 11 and migrating the nucleic acid sample after reaction into another microtube 18.

As a result of the above treatments, a nucleic acid sample composed of mRNAs or cDNAs containing no LEES GENE is recovered in the microtube 18.

FIG. 7 and FIGS. 8A and 8B respectively illustrate apparatus useful for another embodiment of the treatment shown in FIGS. 6D through 6F.

FIG. 7 demonstrates a centrifuge tube having dual-membrane structure as the apparatus. First chamber 701 is equipped with a lid at the top and membrane filter 703 at the bottom. The membrane filter 703 has a configuration for retaining the probe 14 physically or

chemically. As the membrane filter, use can be made of membranes each composed of a structure having sufficient pore size to allow nucleic acids to permeate therethrough, but not to allow nucleic acids hybridized with a probe which is immobilized onto, for example, bead to permeate physically therethrough; or membranes using a substance having affinity to the probe, such as a streptavidin-coated membrane to which biotinylated probe can be bound. The top of second chamber 704 is engaged to the lower part of the first chamber 701, and the chamber 704 is provided with membrane filter 705 in its inside. As the membrane filter 705, use can be made of filters that can retain nucleic acids but allow water and inorganic salts to permeate therethrough to concentrate the nucleic acids, such as conventional ultrafilters. The top of third chamber 706 is engaged to the bottom of the second chamber 704. Centrifuge tubes having analogous structure to that of the above centrifuge tube include, for example, Micropure EZ and Microcon (trade names; Amicon, Danvers, Mass., USA).

Upon the three chambers being assembled, the nucleic acid sample prepared by the aforementioned methods is put into the first chamber 701, and centrifuged by proper centrifugal force; then, the LEES GENE 16 hybridized with the probe which is immobilized onto, for example, beads remains in the first chamber 701, and the rare expressed genes 15 are captured within the second chamber 704. Naturally, LEES GENES which cannot be hybridized with the

probe 14 remain with the rare expressed gene 15 in the second chamber 704. Into the third chamber 706 is collected unnecessary liquid.

5 The genes captured in the second chamber 704 are composed of not only the rare expressed genes 15 but also LEES GENES which cannot be hybridized and treated with the probe 14, and the residual LEES GENES should be treated with another probe. This treatment process will be described later.

10 FIGS. 8A and 8B illustrate a sectional view and a plane view, respectively, of another embodiment in which the isolation of genes is carried out not by centrifuge but by electrophoresis. Electrophoresis cell 801 is provided in its inside with membrane filters 803 and 805.
15 The membrane filter 803 may have the same configuration as the membrane filter 703, whereas the membrane filter 805 may have the same configuration as the membrane filter 705. The inside of the electrophoresis cell 801 constitutes three cells 802, 804 and 806 respectively
20 divided by the membrane filters 803 and 805. The left end of the cell 802 is provided with electrode 811, and the right end of the cell 806 is provided with electrode 812. To each of the electrodes is supplied a given electric potential from power source 810.

25 The nucleic acid sample prepared by the aforementioned method is placed into the cell 802, and a proper buffer is put into the cells 804 and 806. When the amount of the nucleic acid sample in the cell 802 is small,

the buffer should preferably be put into the cell 802. Upon supplying a given electric potential from the power source 810, the LEES GENE 16 hybridized with the probe which is immobilized onto, for example, beads remains in the cell 802, and the rare expressed gene 15 is captured in the cell 804. Naturally, LEES GENES not hybridized with the probe 14 remain with the rare expressed genes 15 in the cell 804.

The genes captured in the cell 804 are composed of not only the rare expressed gene 15 but also LEES GENES which cannot be hybridized with the probe 14, and the residual LEES GENES should be treated with another probe, as in the embodiment shown in FIG. 7.

In the configurations illustrated in FIG. 7 and FIGS. 8A and 8B, the driving force to collect genes to the second chamber 704 or to the cell 804 is not limited to those shown in the above embodiments. The genes can also be collected by transferring the sample solution physically with, for instance, a pump.

Next, the third method will be described in detail with reference to FIG. 9. The third method is a technique to enhance the precision of removal in the first and second methods. It is preferable to remove all LEES GENES by treating once, but it may not be always successful. In order to remove plural LEES GENES with high precision, it is effective to remove the LEES GENES while detecting the removal of individual LEES GENES. According to the third method, therefore, the extent of removal in the recovery

obtained by the first method or second method is monitored, and a successive removing process of genes to be removed is conducted according to necessity.

FIG. 9 illustrates an embodiment for enhancing the precision of the resultant sample obtained by the second method. Step 901 is a step for providing a nucleic acid sample, step 902 is a step for hybridizing probe carriers with LEES GENE and step 903 is a step for the collection of nucleic acid sample containing non-hybridized genes, and these steps are conducted in a similar manner as in the second method.

Step 904 is a step for checking the amount and purity of nucleic acids hybridized with the probe. By taking the embodiment using the beads 13 as illustrated in FIG. 6A through 6F, the step 904 is conducted in the following manner: To the bead cluster 17 shown in FIG. 6E (the bead cluster after the recovery of nucleic acid sample) is added a buffer, and the mixture is heated to denature the hybridized genes. The denatured LEES GENE is quantified by, for example, RT-PCR (reverse transcriptase-PCR, refer for example, FEBS Letters (1994) 231-236). Step 905 is a step for the judgment whether the removing step should be repeated or not, in which the necessity of repeat of the removing step is judged depending on the result of quantification obtained in the step 904.

Conducting this judgment means the removing step should be repeated until the LEES GENE contained in the recovered nucleic acid sample have an equivalent

concentration to that of the rare expressed genes. The judgment can be carried upon the collected non-hybridized nucleic acid sample. When the repetition of the removing step is judged necessary, the steps 902 to 903 are repeated.

5 When it is judged unnecessary, a nucleic acid sample predominantly composed of rare expressed genes is obtained, as illustrated in step 906. To be more specific, once the LEES GENE has an equivalent concentration to those of rare expressed genes, genes which are originally rare expressed
10 genes become major genes occupying sufficient parts of the nucleic acid sample.

By repeating the removing step while quantifying the LEES GENES as described above, the precision of removal can be enhanced. The use of a DNA tip composed of arrayed
15 LEES GENE as the probe carrier instead of beads can provide real-time quantitation of the LEES GENES.

As thus described, gene analysis is conducted on a nucleic acid sample predominantly composed of target rare expressed genes and hence the results of analysis with high
20 precision can be obtained.

The present invention will be further illustrated with reference to the following verification experiment which is not directed to limiting the scope of the invention.

25 Verification Experiment

To verify the effectiveness and advantages of the preparation of a nucleic acid sample predominantly composed of rare expressed genes as in the present

invention, effects of the presence of a LEES GENE on the analysis of rare expressed genes were examined. As an experimental material, ten murine genes having different sizes yet having the same sequences at both ends were used.

5 A series of nucleic acid samples in which the amount of one species of gene (gene H) of ten species (from gene A to gene J in decreasing order of molecular weight) were prepared, and subjected to analysis according to the differential display method using the sequences of the
10 both ends. The analysis will be illustrated in detail below.

The nucleic acid sample was prepared in the following manner. Initially, the ten murine genes were prepared as follows as follows: According to the fluorescent
15 differential display method (Takamichi Muramatsu, Protocols for PCR Experiments of Plants, new edition, 138-143, Shujun-sha, Japan), murine RNAs were isolated on electrophoresis, ten genes, genes A through H having different sizes were cut from gel and cloned to a vector
20 (pGEM-T vector System; Promega, Madison, WS., USA) respectively. Plasmid to be used herein was purified using a commercially available kit (RPM kit; BIO 101 Inc., Calif., USA). Using the purified plasmid as a template, genes A through J were respectively amplified by
25 subjecting to PCR (AmpliTaQ and GeneAmp PCR System 9600; Perkin-Elmer, Norwalk, Conn., USA) according to a conventional method using universal primers (Sequence Listing: Sequence ID Nos. 1 and 2). The PCR product was

purified using QIAquick PCR Purification Kit (QIAGEN Inc., Chatsworth, Calif., USA) and subjected to quantitative analysis with a spectrometer (U-3210, Hitachi Ltd., Japan). Using individual quantified genes, two series of samples, a total of 12 samples, in each series other genes than gene H were constant in amount and the gene H (LEES GENE) was present as much in amount as the other genes by a factor of 1, 100, 500, 1000, 5000 and 10000. Table 1 demonstrates the compositions of the nucleic acid samples.

Table 1: Composition Ratios of the Nucleic Acid Samples (mol/ μ l)

Lane	1	2	3	4	5	6
Ratio of LEES GENE	1	100	500	1000	5000	10000
Other genes	1×10^{-20}	1×10^{-20}	1×10^{-20}	1×10^{-20}	1×10^{-20}	1×10^{-20}
Gene H	1×10^{-20}	1×10^{-18}	5×10^{-18}	1×10^{-17}	5×10^{-17}	1×10^{-16}

Lane	7	8	9	10	11	12
Ratio of LEES GENE	1	100	500	1000	5000	10000
Other genes	1×10^{-18}	1×10^{-18}	1×10^{-18}	1×10^{-18}	1×10^{-18}	1×10^{-18}
Gene H	1×10^{-18}	1×10^{-16}	5×10^{-16}	1×10^{-15}	5×10^{-15}	1×10^{-14}

(Amplification of the Nucleic Acid Samples by PCR)

The primer used for the amplification is as follows:
 anchor primer: Sequence listing: sequence ID No. 3
 Arbitrary primer (OPERON TECHNOLOGY, Inc., Alameda, Calif., USA): Sequence listing: sequence ID NO. 4

The sequences of the above primers were present identically at the both ends of the genes A through J. At the 5' end of the anchor primer was labeled with a fluorescent substance (Texas-Red) for the purpose of

detection.

A reaction mixture for PCR was prepared according to the fluorescent differential display method in the following manner:

5 Table 2 demonstrates the composition of a material mixture.

Table 2: Composition of Material Mixture

2.5 mM dNTP (Nippon Gene Co., Ltd., Japan)	0.4 μ l
10 x GeneTaq Buffer (Nippon Gene Co., Ltd., Japan)	2.0 μ l
anchor primer (1 μ M)	5.0 μ l
Arbitrary primer (10 μ M)	1.0 μ l
GeneTaq (Nippon Gene Co., Ltd., Japan)	0.1 μ l
AmpliTag (Perkin-Elmer)	0.1 μ l
Distilled water	10.4 μ l
Total	19 μ l

10 To 1 μ l each of the nucleic acid samples 1 to 12 was added 19 μ l of the mixture shown in Table 2 to give a series of PCR reaction mixtures (20 μ l). The temperature cycle of PCR was set as follows: The first cycle was composed of 94°C for 3 min, 40°C for 5 min and then 72°C for 5 min in this order, and a subsequent cycle was composed of 94°C
15 for 20 sec, 40°C for 2 min and then 72 °C for 1 min in this order, and this subsequent cycle was repeated a total of 24 times, followed by a reaction at 72°C for 5 min.

(Detection of PCR Products)

20 To 2.0 μ l of the PCR product was added an equivalent volume of a loading buffer (98% formamide, 10 mM EDTA, 0.01% Methyl Violet) and the resultant mixture was treated at 80°C for 2 min to give a sample. The sample was subjected

to electrophoresis on a denatured acrylamide gel (6% Long Ranger (FMC Corp., Rockland, Md., USA), 6.1 M urea, 1.2 xTBE) and then to detection with a fluorescence image scanner (FMBIO, TaKaRa, Japan).

5 FIG. 10 demonstrates the results of above detection, in which lanes 1 through 12 are respectively the results of isolation of the PCR products of the nucleic acid samples 1 through 12 shown in Table 1. Lanes 7 through 12 correspond to the nucleic acid samples containing genes 100 times as much as those of the lanes 1 through 6, respectively. Bands A through J respectively indicate products of the individual genes A thorough J. The numerals 100 through 800 indicate base lengths (bp) of molecular weight markers.

15 The lanes 1, 2 and 7, 8 correspond to the reaction products of the nucleic acid samples containing gene H in an equivalent amount and 100 times as much as the other genes, indicating that all genes were amplified uniformly. The lanes 3, 4, 5, 6 and 9, 10, 11, 12 demonstrate the results of the nucleic acid samples containing gene H 500 20 times or more as much as the other genes. The bands A, B, C, D, E, F, G, I and J in these lanes are apparently thin than those in the lanes 1, 2 and 7, 8, although the nucleic acid samples 1 through 6 and 7 through 12 25 respectively contained equivalent amount of the other genes than gene H, indicating that uniform gene amplification is inhibited.

The above results indicate that the presence of a

gene belonging to the abundant class (gene H) in a nucleic acid sample inhibits the detection of genes belonging to the rare class (genes A, B, C, D, E, F, G, I and J), and that contrary to this, removal of an abundant gene (gene H) from a nucleic acid sample containing largely different amounts of genes enables the detection of rare expressed genes (genes A, B, C, D, E, F, G, I and J).

In comparison between the lanes 1 through 6 and lanes 7 through 12, the results were almost equivalent though the latter nucleic acid samples contained genes 100 times as much as those in the former nucleic acid samples, indicating that even if the absolute amounts of rare expressed genes are increased, the amplification of the genes is inhibited as far as an abundant gene is present in a sample.

The results of the verification experiment above indicate that the removal of an abundant gene from a nucleic acid sample is effective in the expression analysis of rare expressed genes. the advantages of this technique were examined using a detection method with the use of PCR technique in the experiment, whereas similar advantages can presumably be obtained as long as a technique with hybridization of a probe or primer is used for the analysis of expressed amounts.

As described above, genetic diagnoses for studying mutation of genes have been conducted, whereas studies upon the relationship between RNA profiling verifying, in general, the expression styles of mRNAs and diseases have

been launched. It is expected that genetic diagnoses utilizing such RNA profiling will be established in near future. A candidate sample for the RNA profiling is a tissue presenting with a symptom, but the practical
5 preparation of mRNA by subjecting a patient to tissue biopsy is attended with technical difficulties and pain of the patient. Separately, the use of peripheral blood as a sample is advantageous for expected application to prophylaxis diagnosis. In this case, the application of
10 the present invention to a nucleic acid samples obtained from peripheral blood is important and such an expanded application can be realized, as described below.

The peripheral blood contains a large quantity of blood cells such as red blood cells, white blood cells and
15 platelets, and, occasionally, trace amounts of other cells such as metastatic tumor cells. When plural species of cells are present in co-existence, more complicated gene expression is expected, and the problems caused by
20 deference in expression amounts should be solved to obtain accurate information particularly on rare expressed genes. Consequently, the present invention is effective for gene expression analysis for the purpose of genetic diagnosis using peripheral blood. The species and amounts of major
25 blood cells are described in "Molecular Biology of the Cell, 3rd edition, p1164 (Kyoiku-sha K. K., Japan)". Of these blood cells, targets to be analyzed are white blood cells other than red blood cells and platelets both of which have no nuclei. The aforementioned database BodyMap

The invention provides preparation methods of nucleic acid samples for analyses of rare expressed genes, and analyzing methods with high resolution based thereupon.

5 [Sequence Listing]

Sequence ID No.: 1

Sequence length: 23

Sequence type: nucleic acid

Strandedness: single

10 Topology: linear

Species of sequence: synthesized DNA

Sequence: AAAGGGGGAT GTGCTGCAAG GCG

Sequence ID No.: 2

Sequence length: 23

15 Sequence type: nucleic acid

Strandedness: single

Topology: linear

Species of sequence: synthesized DNA

Sequence: GCTTCCGGCT CGTATGTTGT GTG

20 Sequence ID No.: 3

Sequence length: 17

Sequence type: nucleic acid

Strandedness: single

Topology: linear

25 Species of sequence: synthesized DNA

Sequence: GTTTTTTTTTT TTTTTTG

Sequence ID No.: 4

Sequence length: 10

Sequence type: nucleic acid

Strandedness: single

Topology: linear

Species of sequence: synthesized DNA

5 Sequence: GGTCTGGTTG